# This Page Is Inserted by IFW Operations and is not a part of the Official Record

## BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- · TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

## IMAGES ARE BEST AVAILABLE COPY.

As rescanning documents will not correct images, please do not report the images to the Image Problem Mailbox.

THIS PAGE BLANK (USPTO)



#### PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION

#### INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6:		(11) International Publication Number:	WO 95/17507	
C12N 15/11, A61K 31/70, C07H 21/00	A1	(43) International Publication Date:	29 June 1995 (29.06.95)	
(21) International Application Number: PCT/EP (22) International Filing Date: 9 December 1994 (		BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, N		
(30) Priority Data: 93120710.4 23 December 1993 (23.12.9) (34) Countries for which the regional or international application was filed:	3) I DE et	Published With international search report		
(71) Applicant (for all designated States.except US): BI TIK GESELLSCHAFT FÜR BIOMOLEKULAR: NOSTIK MBH [DE/DE]; Carl-Giesecke-Strasse 3, Göttingen (DE).	E DIA	G-		
(72) Inventors; and (75) Inventors; and (75) Inventors Applicants (for US only); BRYSCH, N. [DEDDE]: Am Goldgraben 20, D-37073 Götting SCHLINGENSIEPEN, Kurl-Hermann [DEDDE]: SCHLISCEN Reimar [DEDDE]: Am Goldgraben 13, D-37073 (DE). SCHLINGENSIEPEN, Georg-F. [DE/DE]: Agrabe 20, D-37073 Göttingen (DE).	ten (DI Bovend ISIEPE Götting	er N, en		
(74) Agents: MEYERS, Hans-Wilhelm et al.; Deichmani Hauptbahnhof, D-50667 Köln (DE).	nhaus a	m ·		
			DI HERON EMPERATON	

(54) Title: ANTISENSE NUCLEIC ACIDS FOR THE PREVENTION AND TREATMENT OF DISORDERS IN WHICH EXPRESSION OF C-END PLAYS A ROLE

#### (57) Abstract

The present invention is related to an antisense-nucleic acid or effective derivatives thereof hybridizing with an area of the messenger RNA (mRNA) or the DNA, encoding the pl85<sup>mb.2</sup> receptor (also termed c-erb52, HER2 or neu), a pharmaceutical composition, comprising an antisense nucleic acid or effective derivatives thereof hybridizing with an area of the messenger RNA (mRNA) or the district of the messenger than (mRNA) or the district of the manufacturing of a pharmaceutical composition for the treatment of neoplasms and/or immune diseases and/or diseases involving pathological angiogenesis.



#### FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

ΑT	Austria	GB	United Kingdom	MR	Mauritania
ΑU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	. GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgystan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic	SD	Sudan
CG	Congo		of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SI	Slovenia
CI	Côte d'Ivoire	KZ	Kazakhstan	SK	Slovakia
CM	Cameroon	LI	Liechtenstein	SN	Senegal
CN	China	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TG	Togo
CZ	Czech Republic	LV	Latvia	TJ	Tajikistan
Œ	Germany	MC	Monaco	TT	Trinidad and Tobago
DK	Denmark	MD	Republic of Moldova	UA	Ukraine
ES	Spain	MG	Madagascar	US	United States of Americ
Fī	Finland	ML	Mali	UZ	Uzbekistan
FR	France	MN	Mongolia	VN	Viet Nam
GA	Gabon		-		

WO 95/17507 PCT/EP94/04094

### Antisense nucleic Acids for the prevention and treatment of disorders in which expression of c-erbB plays a role

The present invention is related to an antisense-nucleic acid or effective derivatives thereof hybridizing with an area of the messenger RNA (mRNA) or the DNA, encoding the p185°\*\*b8-2 receptor (also termed c-erbB-2, HER2 or neu), a pharmaceutical composition, comprising an antisense nucleic acid or effective derivatives thereof hybridizing with an area of the messenger RNA (mRNA) or the DNA, encoding the c-erbB-2 receptor as well as the use of said antisense nucleic acids and derivatives thereof for the manufacturing of a pharmaceutical composition for the treatment of neoplasms and/or immune diseases and/or diseases involving pathological anglogenesis.

ErbB-2 is a putative growth factor receptor with an intracellular tyrosine kinase activity that is amplified and/or overexpressed by tumor cells in a variety of neoplasms including breast cancer, lung cancer, esophageal and gastric cancer, bile duct carcinoma, bladder cancer and ovarian cancer.

In breast carcinoma patients, an amplification and overexpression of the c-erbB-2 gene in the tumor tissue has been shown to correlate with a poor clinical prognosis. Overexpession of  $p185^{\text{crbs-2}}$  in non-small-cell lung carcinoma has been shown to impart resistance to a number of chemotherapeutic agents.

WO 93/09788 discloses a method for inhibiting the proliferation of cells which contain an erb B2/neu gene site. The method involves administering a therapeutic dose of an oligonucleotide which is capable of forming a colinear triplex with the promoter region of the erb B2/neu gene.

WO 92/19732 discloses sense and antisense oligonucleotides, namely closed oligonucleotides. These compounds may be used pharmacologically as sense or antisense molecules. It is generally described the therapeutic use of oligonucleotides as sense or antisense agents.

WO 92/13063 discloses a method for effecting expression of growth factors and growth factor receptors in cells or in multicellular animals and methods for testing compounds as effectors of transcription of growth factors and growth factor receptors.

The article "Chemically Modified Oligodeoxynucleotide Analogs as Regulators of Viral and Cellular Gene Expression" in Gene Regulation: Biology of Antisense RNA and DNA discloses in general the use of chemically modified oligonucleotides in the antisense technology.

It is an object of the present invention to provide a compound for the treatment of neoplasms and/or immune diseases and/or diseases involving pathological angiogenesis.

The c-erbB-2 antisense-oligonucleotide of the invention solving the problem addressed above have the sequences as disclosed in the sequence listing under Seq. ID No. 1-105, having a DNA- or RNA-type structure. The control oligonucleotide has the sequence as disclosed in the sequence

listing under Seq. ID No 106, having a DNA- or RNA-type structure.

The antisense nucleic acids of the invention, were able to strongly inhibit the expression of the p185\*erb\*-? protein, tyrosine kinase activity and cell growth in a variety of tumor cells including breast cancer cells. Untransformed normal fibroblasts were not growth inhibited by the anti-c-erb\*-2 antisense compounds. This suggests that p185\*erb\*-? plays a pathogenetic role in the growth of the above mentioned tumor cells.

Furthermore, surprisingly, the immune response to a variety of neoplasms was significantly increased by the use of the antisense nucleic acids of the invention. Immune cell growth and activity was stimulated in co-culture assays culturing tumor cells and peripheral blood monocytes together.

Surprisingly, the antisense nucleic acids of the invention, also acted as strong inhibitors of angiogenesis. This suggests, that either the secreted truncated form of the c-erbB-2 protein or the full receptor protein may play a causal role in pathological neoangiogenesis.

According to the invention antisense nucleic acids or effective derivatives thereof which hybridize with an area of the mRNA or DNA coding for p185erb8-2 can effectively treat the diseases addressed above. The antisense nucleic acid is able to hybridize with regions of p185erb8-2 mRNA. It is understood by the skilled person that fragments of the antisense nucleic acids and antisense nucleic acids containing these sequences work according to the invention so long as production of p185erb8-2 is reduced or inhibited.

According to the invention the antisense-oligonucleotides are obtainable by solid phase synthesis using phosphite triester chemistry by growing the nucleotide chain in 3'-5'

direction in that the respective nucleotide is coupled to the first nucleotide which is covalently attached to the solid phase comprising the steps of

- cleaving 5'DMT protecting group of the previous nucleotide,
- adding the respective nucleotide for chain propagation,
- modifying the phosphite group subsequently cap unreacted 5'-hydroxyl groups and
- cleaving the oligonucleotide from the solid support,
- followed by working up the synthesis product.

The chemical structures of oligodeoxy-ribonucleotides are given in figure 1 as well as the respective structures of antisense oligo-ribonucleotides are given in figure 2. The oligonucleotide chain is to be understood as a detail out of a longer nucleotide chain.

In figure 1, lit. B means an organic base such as adenine (A), guanine (G), cytosine (C) and thymine (T) which are coupled via N9(A,G) or N1(D,T) to the desoxyribose. The sequence of the bases is the reverse complement of the genetic target sequence (mRNA-sequence). The modifications used are

1. Oligodeoxy-ribonucleotides where all  $\ensuremath{\text{R}}^1$  are substituted by

1.1 
$$R^1 = 0$$

1.2 
$$R^1 = S$$

1.3 
$$R^1 = F$$

1.4 
$$R^1 = CH_3$$

1.5  $R^1 = OEt$ 

 Oligodeoxy-ribonucleotides where R<sup>1</sup> is varied at the internucleotide phosphates within one oligonucleotide

where B = deoxy-ribonucleotide dA, dC, dG or dT depending

on gene sequence

p = internucleotide phosphate

n = an oligodeoxy-ribonucleotide stretch of length 6 - 20 bases

2.1 
$$R^{1a} = S;$$
  $R^{1b} = 0$   
2.2  $R^{1a} = CH_3;$   $R^{1b} = 0$   
2.3  $R^{1a} = S;$   $R^{1b} = CH_3;$   
2.4  $R^{1a} = CH_3;$   $R^{1b} = S$ 

3. Oligodeoxy-ribonucleotides where  $\mathbb{R}^1$  is alternated at the internucleotide phosphates within one oligonucleotide

where B = deoxy-ribonucleotide dA, dC, dG or dT
 depending on gene sequence

p = internucleotide phosphate

n = an oligodeoxy-ribodinucleotide stretch of length
4 - 12 dinucleotides

3.2 
$$R^{la} = S;$$
  $R^{lb} = 0$   
3.2  $R^{la} = CH_3;$   $R^{lb} = 0$   
3.3  $R^{la} = S;$   $R^{lb} = CH_3$ 

- 4. Any of the compounds 1.1 1.5; 2.1 2.4; 3.1 3.3 coupled at  $\mathbb{R}^2$  with the following compounds which are covalently coupled to increased cellular uptake
- 4.1 cholesterol
- 4.2 poly(L)lysine
- 4.3 transferrin
- 5. Any of the compounds 1.1 1.5; 2.1 2.4; 3.1 3.3 coupled at R<sup>3</sup> with the following compounds which are covalently coupled to increase cellular uptake
- 5.1 cholesterol
- 5.2 poly(L)lysine
- 5.3 transferrin

In the case of the RNA-oligonucleotides (figure 2) are the basis (adenine (A), guanine (G), cytosine (C), uracil (U)) coupled via N9 (A,G) or N1 (C,U) to the ribose. The sequence of the basis is the reverse complement of the genetic target sequence (mRNA-sequence). The modifications in the oligonucleotide sequence used are as follows

- 6. Oligo-ribonucleotides where all R<sup>1</sup> are substituted by
- 6.1  $R^1 = 0$
- 6.2  $R^1 = S$
- $R^1 = F$
- 6.4 R1 = CH3

Oligo-ribonucleotides where  $\mathbb{R}^1$  is varied at the internucleotide phosphates within one oligonucleotide

where B = ribonucleotide A, C, G or T depending on gene sequence

- p = internucleotide phosphate
- n = an oligo-ribonucleotide stretch of length 4 20 bases
- 7.1  $R^{1a} = S;$   $R^{1b} = O$ 7.2  $R^{1a} = CH_3;$   $R^{1b} = O$ 7.3  $R^{1a} = S;$   $R^{1b} = CH_3;$ 7.4  $R^{1a} = CH_3;$   $R^{1b} = S$
- Oligo-ribonucleotides where R<sup>1</sup> is alternated at the internucleotide phosphates within one oligonucleotide

where B = ribonucleotide A, C, G or T depending
 on gene sequence

- p = internucleotide phosphate
- n = an oligo-ribodinucleotide stretch of length 4 -12 dinucleotides
- 8.2  $R^{1a} = S;$   $R^{1D} = O$ 8.2  $R^{1a} = CH_3;$   $R^{1b} = O$ 8.3  $R^{1a} = S;$   $R^{1b} = CH_3$
- 9. Any of the compounds 6.1 6.5; 7.1 7.4; 8.1 8.3 coupled at R<sup>2</sup> with the following compounds which are covalently coupled to increase cellular uptake
- 9.1 cholesterol
- 9.2 poly(L)lysine
- 9.3 transferrin
- 10. Any of the compounds 6.1 6.5; 7.1 7.4; 8.1 8.3

- 8 -

coupled at R3 the following compounds are covalently coupled to increased cellular uptake

- 10.1 cholesterol
- 10.2 poly(L)lysine
- 10.3 transferrin
- 11. Any of the compounds 6.1 6.5; 7.1 7.4; 8.1 8.3; 9.1 9.3; 10.1 10.3 where all  $R^4$  are substituted by
- 11.1  $R^4 = 0$
- 11.2  $R^4 = F$
- 11.3  $R^4 = CH_2$

In a preferred embodiment of the oligonucleotides of the invention they are phosphorothioate derivatives, having a DNAor RNA-type structure.

It is possible that one single individual sequence as mentioned above works as an antisense nucleic acid or oligonucleotide structure according to the invention. However, it is also possible that one strand of nucleotides comprises more than one of the sequences as mentioned above directly covalently linked or with other nucleotides covalently linked in between. Preferably, individual oligonucleotides are addressed.

In a preferred embodiment of these oligo-nucleotides they are phosphorothioate derivatives.

Modifications of the antisense-oligonucleotides are advantageous since they are not as fast destroyed by endogenous factors when applied as this is valid for naturally occurring nucleotide sequences. However, it is understood by the skilled person that also naturally occurring nucleotides having the disclosed sequence can be used according to

the invention. In a very preferred embodiment the modification is a phosphorothioate modification.

The synthesis of the oligodeoxy-nucleotide of the invention is described as an example in a greater detail as follows.

Oligodeoxy-nucleotides were synthesized by stepwise 5'-addition of protected nucleosides using phosphite triester chemistry. The nucleotide A was introduced as 5'dimethoxy-trityl-deoxyadenosine(N-benzoyl)-N,N'-diisopropyl-2-cyanoethyl phosphoramidite (0.1 M); C was introduced by a 5'dimethoxytrityl-deoxycytidine(N<sup>4</sup>-benzoyl)-N,N'-diisopropyl-2-cyanoethyl phosphoramidite; G was introduced as 5'-dimethoxytrityl-deoxyguanosine(N<sup>8</sup>-isobutyryl)-N,N'-diisopropyl-2-cyanoethyl phosphoramidite and the T was introduced as 5'-dimethodytrityl-deoxythymidine-N,N'-diisopropyl-2-cyanoethyl phosphoramidite. The nucleosides were preferably applied in 0.1 M concentration dissolved in acetonitrile.

Synthesis was performed on controlled pore glass particles of approximately 150  $\mu m$  diameter (pore diameter 500 Å) to which the most 3' nucleoside is covalently attached via a long-chain alkylamine linker (average loading 30  $\mu mol/g$  solid support).

The solid support was loaded into a cylindrical synthesis column, capped on both ends with filters which permit adequate flow of reagents but hold back the solid synthesis support. Reagents were delivered and withdrawn from the synthesis column using positive pressure of inert gas. The nucleotides were added to the growing oligonucleotide chain in 3'-> 5' direction. Each nucleotide was coupled using one round of the following synthesis cycle:

Cleave 5'DMT (dimethoxytrityl) protecting group of the previous nucleotide with 3-chloroacetic acid in dichloromethane followed by washing the column with anhydrous acetonitrile.

1

Then simultaneously one of the bases in form of their protected derivative depending on the sequence was added plus tetrazole in acetonitrile. After reaction the reaction mixture has been withdrawn and the phosphite was oxidized with a mixture of sulfur  $(S_8)$  in carbon disulfide/pyridine/triethylamine. After the oxidation reaction the mixture was withdrawn and the column was washed with acetonitrile. The unreacted 5'-hydroxyl groups were capped with simultaneous addition of 1-methylimidazole and acetic anhydride/lutidine/tetrahydrofuran. Thereafter, the synthesis column was washed with acetonitrile and the next cycle was started.

The work up procedure and purification of the synthesis products occurred as follows.

After the addition of the last nucleotide the deoxynucleotides were cleaved from the solid support by incubation in ammonia solution. Exocyclic base protecting groups were removed by further incubation in ammonia. Then the ammonia was evaporated under vacuum. Full-length synthesis products still bearing the 5'DMT protecting group were separated from shorter failure contaminants using reverse phase high performance liquid chromatography on silica C, stationary phase. Eluents from the product peak were collected, dried under vacuum and the 5'-DMT protecting group cleaved by incubation in acetic acid which was evaporated thereafter under vacuum. The synthesis products were solubilized in the deionized water and extracted three times with diethylether. Then the products were dried in vacuo. Another HPLC-AX chromatography was performed and the eluents from the product peak were dialyzed against excess of Trisbuffer as well as a second dialysis against deionized water. The final products were lyophilized and stored dry.

The antisense-nucleic acid of the invention can be used as pharmaceutical composition or medicament. This medicament can be used for treating neoplasms and/or immune diseases and/or diseases involving pathological angiogenesis in which the expression of c-erbB-2 derived receptor protein or truncated p185c-erbB2 is of relevance for the pathogenicity. It can be used to reduce neoplastic cell growth in cells expressing p185c-erbB2, to reverse resistance of tumor cells to the immune-response, to inhibit pathological angiogenesis and to stimulate the immune system.

The antisense nucleic acids of the invention are intermediate products of the pharmaceutical composition or medicament of the invention. The pharmaceutical composition may comprise besides the effective compound(s) suitable carrier agents, solvents and other ingredients known in the art for producing medicaments. Preferably, these agents facilitate the administration of the pharmaceutical composition of the invention. Typically, the pharmaceutical composition is administered as i.v. infusion or i.v. bolus injection. The amount of the active ingredient to be administered is typically in the range of 0.2 - 50 mg of the oligonucleotide per kg body weight per day, in particular 1 - 12 mg/kg body weight per day.

In principal the compound which can be used as an active compound in the pharmaceutical composition can be used as a diagnostic tool for evaluating whether the respective genes are expresses. Typically, radio active labelled nucleotides are hybridized by the method of northern blotting which is well-known in the art or in situ with a sample to be examined. The degree of hybridization is a measure for the degree of expression of the respective genes.

The effect of c-erbB2 specific antisense-oligonucleotides on neoplastic cell growth was investigated. It was demonstrated that antisense oligodeoxynucleotides as well as phosphorothioate modified nucleic acids, complementary to c-erbB2 mRNA could specifically inhibit pl85<sup>c-erbB2</sup> protein expression and could to a significant amount reduce cell

proliferation in breast cancer cells, ovarian carcinoma cells and bladder cancer cells. Also, it could be shown that protein synthesis and S6 kinase activity were strongly reduced in tumor cells, treated with the antisense nucleic acid.

Furthermore, the immune response to a variety of neoplasms was significantly increased by the use of the antisense nucleic acids described below. Lymphocyte growth and activity was stimulated in co-culture assays culturing tumor cells and peripheral blood monocytes together.

Furthermore, the antisense nucleic acids described above, also acted as inhibitors of angiogenesis.

- 13 -

#### SEQUENCE LISTING

- (1) GENERAL INFORMATION:
  - (i) APPLICANT:
    - (A) NAME: Biognostik Gesellschaft fuer biomolekulare Diagnostik mbH
      - (B) STREET: Carl-Giesecke-Str. 3
      - (C) CITY: Goettingen
      - (E) COUNTRY: Germany
      - (F) POSTAL CODE (ZIP): 37079
  - (ii) TITLE OF INVENTION: Antisense nucleic Acids for the prevention and treatment of disorders in which expression of c-erbB plays a role
    - (iii) NUMBER OF SEQUENCES: 106
    - (iv) COMPUTER READABLE FORM:
      - (A) MEDIUM TYPE: Floppy disc
      - (B) COMPUTER: IBM PC compatible
      - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
      - (D) SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
    - (v) CURRENT APPLICATION DATA: APPLICATION NUMBER: EP 93120710.4
    - (2) INFORMATION FOR SEQUENCE ID NO: 1:
      - (i) SEQUENCE CHARACTERISTICS:
        - (A) LENGTH: 14 base pairs
        - (B) TYPE: nucleic acid
        - (C) STRANDEDNESS: unknown
        - (D) TOPOLOGY: unknown
      - (ii) MOLECULE TYPE: DNA (genomic)
      - (iii) ANTI-SENSE: YES
      - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

TTCATGTCTG TGCC

14

- (2) INFORMATION FOR SEQUENCE ID NO: 2:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 14 base pairs
    - (B) TYPE: nucleic acid
    - (C) STRANDEDNESS: unknown
      (D) TOPOLOGY: unknown
    - (ii) MOLECULE TYPE: DNA (genomic)
    - (iii) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:	
GTAGGTGAGT TCCA	14
(2) INFORMATION FOR SEQUENCE ID NO: 3:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEONESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:	
GTTGTGAGCG ATGA	14
(2) INFORMATION FOR SEQUENCE ID NO: 4:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 18 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:	
CATAGTTGTC CTCAAAGA	18
(2) INFORMATION FOR SEQUENCE ID NO: 5:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:	
GGCATAGTTG TCCT	

(2) INFORMATION FOR SEQUENCE ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:	
CATTGTCTAG CACG	14
(2) INFORMATION FOR SEQUENCE ID NO: 7:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:	
CTCCATTGTC TAGC	14
(2) INFORMATION FOR SEQUENCE ID NO: 8:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:	
GTATTGTTCA GCGG	14
* .	

(2) 114	FORMATION TOR SEQUENCE ID NO. 3.	
(	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(	(ii) MOLECULE TYPE: DNA (genomic)	
(	(iii) ANTI-SENSE: YES	
(	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:	
TCAAGA	ATCTC TGTGAG	16
(2) IN	FORMATION FOR SEQUENCE ID NO: 10:	
(	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(	(ii) MOLECULE TYPE: DNA (genomic)	
(	(iii) ANTI-SENSE: YES	
	(Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:	16
		10
(2) IN	NFORMATION FOR SEQUENCE ID NO: 11:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:	
TCCTTC	CCACA AAATCG	16
		-

(2) INFORMATION FOR SEQUENCE ID NO: 12:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
÷
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:
GTGGAAGATG TCCT 1
(2) INFORMATION FOR SEQUENCE ID NO: 13:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:
TCTTGTGGAA GATGTC 1
(2) INFORMATION FOR SEQUENCE ID NO: 14:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:
TCTATCAGTG TGAGAG 1

(2) INFORMATION FOR SEQUENCE ID NO: 15:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:	
GGTTGGTGTC TATC	14
(2) INFORMATION FOR SEQUENCE ID NO: 16:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:	
ACATCGGAGA ACAG	14
(2) INFORMATION FOR SEQUENCE ID NO: 17:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:	(
CCTTACACAT CGGA	14
and.	

**	
(2) INFORMATION FOR SEQUENCE ID NO: 18:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:	
CAATCCTCA GAACTC	16
(2) INFORMATION FOR SEQUENCE ID NO: 19:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:	
GCTCTGACAA TCCT	14
(2) INFORMATION FOR SEQUENCE ID NO: 20:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:	
TGGTTGAAGT GGAG	14

(2) INFORMATION FOR SEQUENCE ID NO: 21:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:	
CTGTGGTTGA AGTG	4
(2) INFORMATION FOR SEQUENCE ID NO: 22:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:	
GTTGTAGGTG ACCA 1	4
(2) INFORMATION FOR SEQUENCE ID NO: 23:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:	
CTGTGTTGTA GGTG	4

(2)	INFORMATION FOR SEQUENCE ID NO: 24:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:	
GACT	CAAACG TGTC	
(2)	INFORMATION FOR SEQUENCE ID NO: 25:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:	
CATG	GACTCA AACG	
(2)	INFORMATION FOR SEQUENCE ID NO: 26:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
<i>(</i> -	(iii) ANTI-SENSE: YES	
/	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:	

(2) INFORMATION FOR SEQUENCE ID NO: 27:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:	
CCGAATGTAT ACCG	14
(2) INFORMATION FOR SEQUENCE ID NO: 28:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:	
GCCGAATGTA TACC .	14
(2) INFORMATION FOR SEQUENCE ID NO: 29:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYFE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:	
GTAGTTGTAG GGAC	14

WO 95/17507

#### PCT/EP94/04094

- 23 -

(2)	INFORMATION FOR SEQUENCE ID NO: 30:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 18 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:	
TAGA	AAGGTA GTTGTAGG	18
(2)	INFORMATION FOR SEQUENCE ID NO: 31:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:	
GTAG	AAAGGT AGTTGTAG	18
(2)	INFORMATION FOR SEQUENCE ID NO: 32:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDENDESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 32:	
CGTA	AGAAAGG TAGTTG	16

(2) INFORMATION FOR SEQUENCE ID NO: 33:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 33:	
CCGTAGAAAG GTAG	14
(2) INFORMATION FOR SEQUENCE ID NO: 34:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:	
GACCATAGCA CACT	14
(2) INFORMATION FOR SEQUENCE ID NO: 35:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35:	
GGATATTGGC ACTG	14

(2) INFORMATION FOR SEQUENCE ID NO: 36:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:	
CCTGGATATT GGCA	14
(2) INFORMATION FOR SEQUENCE ID NO: 37:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 37:	
GCTCCCAAAG ATCT	14
(2) INFORMATION FOR SEQUENCE ID NO: 38:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 38: CCCATCAAAG CTCT	1

(2) INFORMATION FOR SEQUENCE ID NO: 39:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 39:	
CAAACACTTG GAGC	14
(2) INFORMATION FOR SEQUENCE ID NO: 40:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:	
GTCTCAAACA CTTGGA	16
(2) INFORMATION FOR SEQUENCE ID NO: 41	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 41:	
GAGTCTCAAA CACTTG	16

(2) INFORMATION FOR SEQUENCE ID NO: 42:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:	
GTAACCTGTG ATCTCT	16
(2) INFORMATION FOR SEQUENCE ID NO: 43:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDENDESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 43:	
GGTAACCTGT GATC	14
(2) INFORMATION FOR SEQUENCE ID NO: 44:	*
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 44:	
GTATAGGTAA CCTGTG	16

(2) INFORMATION FOR SEQUENCE ID NO: 45:
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 18 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45:
TGAGATGTAT AGGTAACC
(2) INFORMATION FOR SEQUENCE ID NO: 46:
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:
TGCTGAGATG TATAGG
(2) INFORMATION FOR SEQUENCE ID NO: 47:
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 47:
CCATGCTGAG ATGT

(2) INFORMATION FOR SEQUENCE ID NO: 48:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 48:	
GGATTACTTG CAGG	1
(2) INFORMATION FOR SEQUENCE ID NO:49:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:	
TGTTATGGTG GATGAG	1
(2) INFORMATION FOR SEQUENCE ID NO: 50:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs (b) TYPE: nucleic acid (c) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 50:	
GGTGTTATGG TGGA	1

(1) INFORMATION FOR BEGGENCE ID NO. 31:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 51:	
GCAGTTGACA CACT	14
(2) INFORMATION FOR SEQUENCE ID NO: 52:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 52:	
AGTACTCGGC ATTC	14
(2) INFORMATION FOR SEQUENCE ID NO: 53:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 53:	
CATTCACATA CTCCCT	16

(2) INFORMATION FOR SEQUENCE ID NO: 54:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54:
TCCAAAACAG GTCACT . 10
(2) INFORMATION FOR SEQUENCE ID NO: 55:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 55:
GGTCCTTATA GTGG
(2) INFORMATION FOR SEQUENCE ID NO: 56:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 56:
CAGAATGCCA ACCA 1

(2) INFORMATION FOR SEQUENCE ID NO: 57:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 57:	14
ACGAGAATGC CAAC	14
(2) INFORMATION FOR SEQUENCE ID NO: 58:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:	
GATCCCAAAG ACCA	14
(2) INFORMATION FOR SEQUENCE ID NO: 59:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:	
TCGCTTGATG AGGA	14

(2) INFORMATION FOR SEQUENCE ID NO: 60:

	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
1	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:	
CATO	CGTGTAC TTCC	1
(2)	INFORMATION FOR SEQUENCE ID NO: 61:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:	
GCA:	TCGTGTA CTTC	. 1
(2)	INFORMATION FOR SEQUENCE ID NO: 62:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: unknown (D) TOPOLOGY: unknown	-
	(ii) MOLECULE TYPE: DNA (genomic)	
/	(iii) ANTI-SENSE: YES	
ACT	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:	1
,	<b>V</b>	

(2) INFORMATION FOR SEQUENCE ID NO: 63:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 63:
CTTGTAGACT GTGC 14
(2) INFORMATION FOR SEQUENCE ID NO: 64:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 64:
CCCTTGTAGA CTGT 14
(2) INFORMATION FOR SEQUENCE ID NO: 65:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 65:
TCAACACTTT GATGGC 16

(2) INFORMATION FOR SEQUENCE ID NO: 66:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:
CCCTCAACAC TTTG
(2) INFORMATION FOR SEQUENCE ID NO: 67:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 67:
GTGTTTTCCC TCAACA 10
(2) INFORMATION FOR SEQUENCE ID NO: 68:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 68:
GTATGCTTCG TCTAAG 1

(2) INFORMATION FOR SEQUENCE ID NO: 69:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 69:	
CGTATGCTTC GTCT	14
(2) INFORMATION FOR SEQUENCE ID NO: 70:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70:	
CCATCACGTA TGCT	14
(2) INFORMATION FOR SEQUENCE ID NO: 71:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 71:	
GCATAAGCTG TGTC	1.4

(2) INFORMATION FOR SEQUENCE ID NO: 72:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:	
CATGGTCTAA GAGG	14
(2) INFORMATION FOR SEQUENCE ID NO: 73:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:	
CAATCTGCAT ACACCA	16
(2) INFORMATION FOR SEQUENCE ID NO: 74:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74:	
GGCAATCTGC ATAC	14

(2) INFORMATION FOR SEQUENCE ID NO: 75:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 75:
CTGTCTCGTC AATG
(2) INFORMATION FOR SEQUENCE ID NO: 76:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 76:
CATAACTCCA CACATC 16
(2) INFORMATION FOR SEQUENCE ID NO: 77:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown
(ii) MOLECULE TYPE: DNA (genomic)
(iii) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 77:
AGTCACACCA TAACTC

(2) INFORMATION FOR SEQUENCE ID NO: 78:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 78:	
ACAGTCACAC CATAAC	1
(2) INFORMATION FOR SEQUENCE ID NO: 79:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
· · · · · · · · · · · · · · · · · · ·	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79:	
CCCCAAAAGT CATC ·	1
(2) INFORMATION FOR SEQUENCE ID NO: 80:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80: TCGTAAGGTT TGGC	1

(2) II	NFORMATION FOR SEQUENCE ID NO: 81:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 81:	
GATCCC	CATCG TAAG	14
(2) IN	NFORMATION FOR SEQUENCE ID NO: 82:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 82:	
CAATGG	GTGCA GATG	14
(2) IN	NFORMATION FOR SEQUENCE ID NO: 83:	
1	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
	(ii) MOLECULE TYPE: DNA (genomic)	
1	(iii) ANTI-SENSE: YES	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83:	
	CAATG GTGC	
		14

2) INFORMATION FOR SEQUENCE ID NO: 84:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 84:	
STAGACATCA ATGGTG	16
(2) INFORMATION FOR SEQUENCE ID NO: 85:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 18 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 85:	
CATGATCATG TAGACATC	18
(2) INFORMATION FOR SEQUENCE ID NO: 86:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 86:	
CCATGATCAT GTAGAC	16

(2) INFORMATION FOR SEQUENCE ID NO: 87:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 18 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDMESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 87:	
CATTTGACCA TGATCATG	18
(2) INFORMATION FOR SEQUENCE ID NO: 88:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: unknown  (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 88:	
CCAACATTTG ACCATG	16
(2) INFORMATION FOR SEQUENCE ID NO: 89:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 base pairs	
(B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(E) TYPE: nucleic acid (C) STRANDEDNESS: unknown	
(B) TYPE: nucleic acid (C) STRANDEDMESS: unknown (D) TOPOLOGY: unknown	
(B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown (ii) MOLECULE TYPE: DNA (genomic)	
(B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown (ii) MOLECULE TYPE: DNA (genomic)	
(B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown (ii) MOLECULE TYPE: DNA (genomic) (iii) ANTI-SENSE: YES	18

- 43	
(2) INFORMATION FOR SEQUENCE ID NO: 90:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 90:	
GAGTCAATCA TCCAACAT	18
(2) INFORMATION FOR SEQUENCE ID NO: 91:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 91:	
CAGAGTCAAT CATCCA	16
(2) INFORMATION FOR SEQUENCE ID NO: 92:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 92:	
CCGACATTCA GAGT	14

(2) INFORMATION FOR SEQUENCE ID NO: 93:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 93:	
	1
(2) INFORMATION FOR SEQUENCE ID NO: 94:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 94:	
	14
(2) INFORMATION FOR SEQUENCE ID NO: 95:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 95:	
CCATCAAATA CATCGG	16

(2) INFORMATION FOR SEQUENCE ID NO: 96:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 96:	
TCACCATCAA ATACATCG	18
(2) INFORMATION FOR SEQUENCE ID NO: 97:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 97:	
CAACGTAGCC ATCA	14
(2) INFORMATION FOR SEQUENCE ID NO: 98:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 98:	
ACGTCTTTGA CGAC	1

(2) INFORMATION FOR SEQUENCE ID NO: 99:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 99:	
CAAAAACGTC TTTGACGA	L 8
(2) INFORMATION FOR SEQUENCE ID NO: 100:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 100:	
GGCAAAAACG TCTTTG	L6
(2) INFORMATION FOR SEQUENCE ID NO: 101:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 101:	
CAAAGGCAAA AACGTC	16

(2) INFORMATION FOR SEQUENCE ID NO: 102:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 102:	
GTGTCAAGTA CTCG	14
(2) INFORMATION FOR SEQUENCE ID NO: 103:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 103:	
STAATAGAGG TTGTCG	1
(2) INFORMATION FOR SEQUENCE ID NO: 104:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:	
CCCAOTANIA GAOG	1.

(2) INFORMATION FOR SEQUENCE ID NO: 105:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 105:	
CATGGTGCTC ACTG	14
(2) INFORMATION FOR SEQUENCE ID NO: 106:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGH: 14 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: unknown (D) TOPOLOGY: unknown	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 106:	

## Claims

- An antisense nucleic acid or an effective derivative therefrom which is capable of treating or preventing neoplasms, immune diseases and/or diseases involving pathological angiogenesis, hybridizing with an area of the messenger RNA (mRNA) and/or DNA encoding c-erbB-2, comprising the following sequences identified in the listing under Seq. ID No. 1 - 105, having DNA- or RNA-type structure.
- Antisense oligonucleotides of claim 1 wherein the oligonucleotides are modified oligonucleotides such as phosphorothioate derivatives.
- 3. Antisense nucleic acid or -oligonucleotides according to any one of the claims 1 and/or 2 obtainable by solid phase synthesis using phosphite triester chemistry by growing the nucleotide chain in 3'-5' direction in that the respective nucleotide is coupled to the first nucleotide which is covalently attached to the solid phase comprising the steps of
  - cleaving 5'DMT protecting group of the previous nucleotide,
  - adding the respective nucleotide for chain propagation.
  - modifying phosphite groups subsequently cap unreacted 5'-hydroxyl groups and
  - cleaving the oligonucleotide from the solid support,
  - followed by working up the synthesis product.

- 4. A pharmaceutical composition comprising an effective amount of a compound of any one of the claims 1 to 3 for the prevention and treatment of neoplasms and/or immune diseases and/or diseases involving pathological angiogenesis.
- 5. Use of a compound according to any one of the claims 1 to 3 for the preparation of a pharmaceutical composition for the prevention and treatment of neoplasms and/or immune diseases and/or diseases involving pathological angiogenesis.
- Use of a compound according to any one of the claims
   to 3 as diagnostic agent.
- 7. Method of treating or preventing neoplasms and/or immune diseases and diseases involving pathological angiogenesis by administering an effective amount of the compound according to any one of the claims 1 to 3 or a pharmaceutical composition of claim 4 to a patient suffering from disorders related with the expression of c-erbB-2

		PCT/EP 94	/04094
A. CLASSI IPC 6	FICATION OF SUBJECT MATTER C12N15/11 A61K31/70 C07H21/00	)	
	o International Patent Classification (IPC) or to both national classific	ation and IPC	
	ocumentation searched (classification system followed by classification	n symbols)	
IPC 6	C12N A61K		
Documentat	ion searched other than minimum documentation to the extent that su	eh documents are included in the fields s	earched
Electronie d	ata base consulted during the international search (name of data base	and, where practical, search terms used)	
C. DOCUM	IENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the rel-	evant passages	Relevant to claim No.
Y	ERICKSON, R. & IZANT, J. 'Gene re- biology of antisense RNA and DNA' RAVEN PRESS, Ltd., NEW YORK, USA pages 317-328. SCHLINGENSIEPEN, KH- & BRYSCH, ' 'Phosphorothicate oligomers: inhi oncogene expression in tumor cells and too gene function analysis' see the whole document	W.: bitors of 1s for	1-7
	ther documents are listed in the continuation of box C.	X Patent family members are liste	in annex.
'A' docur consi 'E' earliet filing 'L' docur which citath 'O' docur	when defining the general state of the art which is not derived to be of particular relevance of comment but published on or after the international data wheth may shrow counter on referred dustrief or rest wheth may shrow counter our referred dustrief or one other present the southern one state of another one other present internation (as appendixed) ment referrent to an oral disclosure, use, exhibition or meant with the counter of the present of the present of the present of the present present of the present present present of the present pr	Take roomment published after the iter princip date and not in notice or princip date and not in notice or princip date and not in notice or princip date and not in order or invention.  "A document of particular retevance; if cannot be considered novel or can review as inventive as the order of comment of particular retevance; if document it combined with one or ments, and combination being of met ant.  "As document member of the same pate."	with the application of theory underlying the the claimed invention to be considered to document is taken alone the claimed invention inventive step when the more other such docu- tous to a person skilled
	e actual completion of the international search	Date of mailing of the international	search report
	28 March 1995	0 4 -04- 1	995

Form PCT/ISA/210 (second sheet) (July 1992)

Name and mailing address of the ISA

ing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Face (+31-70) 340-3016

Authorized officer

Andres, S

. 1

C/Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. 1,2,4-7 SCIENCE. vol. 230. 6 December 1985 LANCASTER, PA US. pages 1132-1139. COUSSENS, L. ET AL. 'Tyrosine kinase receptor with extensive homology to EGF receptor shares chromosomal location with neu oncogene' see figure 3 WO.A.92 19732 (GENSET) 12 November 1992 3 see page 22, line 0 - page 23, line 1 see page 32, line 26 - page 33, line 9 1-7 PROCEEDINGS OF THE AMERICAN ASSOCIATION 0.A FOR CANCER RESEARCH, vol. 32, March 1991 page 433 BRYSCH, W. ET AL. 'Inhibiting c-erbB-2 overexpression in human mammary carcinoma cells with phosphorothicate oligodeoxynucleotides' see abstract 1,4-7 WO.A.93 09788 (BAYLOR COLLEGE OF MEDICINE) A 27 May 1993 see page 2, line 19 - page 3, line 10 see page 7, line 3 - line 16 see claims 1.4-7 WO.A.92 13063 (ONCOGENE SCIENCE, INC.) 6 August 1992 August 1992 see page 5, line 21 - line 35 see page 15, line 23 - page 16, line 6 see page 31, line 18 - page 32, line 15 see page 49, line 14 - page 50, line 12 see claims 2,15-20,66,111-115 see claims 127-133

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

1

## INTERNATIONAL SEARCH REPORT

Int ational application No.
PCT/EP 94/04094

D	
Box I	Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
This inte	rnational search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. <b>X</b>	Claims Non:: because they relate to subject matter not required to be searched by this Authority, namely: Remark: Although claim 7 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. 🗌	Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. 🗌	Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II	Observations where unity of invention is lacking (Continuation of item 2 of first sheet)
This Inte	raational Searching Authority found multiple inventions in this international application, as follows:
ı. 🔲	As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2.	As all searchable claims could be searches without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
з. 🗌	As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos:
4.	No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims, it is covered by claims Nos.:
Remark e	The additional search fees were accompanied by the applicant's protest.  No protest accompanied the payment of additional search fees.

Patent document	Publication	Patent	family	Publication
WO-A-9219732	12-11-92	FR-A- AU-A- CA-A- EP-A- JP-T-	2675803 1759692 2102229 0581848 6506834	30-10-92 21-12-92 26-10-92 09-02-94 04-08-94
WO-A-9309788	27-05-93	NONE		
WO-A-9213063	06-08-92	AU-A-	1469292	27-08-92
1-				
i.				
-1				
				•
•				
/				
/				
	•			
/				